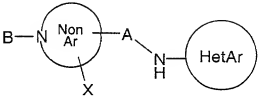
WHAT IS CLAIMED IS:

1. A compound having the formula (I):



(I)

or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 5-7 membered ring containing 1 or 2 nitrogen ring atoms or an aza bicyclo octane ring;

HetAr is a 5 or 6 membered heteroaromatic ring containing 1-3 nitrogen ring atoms, or isoxazolyl, thiazolyl, thiadiazolyl, quinolinyl, quinazolinyl, purinyl, pteridinyl, benzimidazolyl, pyrrolopyrimidinyl, or imidazopyridinyl;

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)HNCH_2–, Si$(CH_3)_3$–C–, or $NH_2C(O)$–;

A is –C_{0-4}alkyl–;

B is aryl$(CH_2)_{0-3}$–O–C(O)–, heteroaryl$(CH_2)_{1-3}$–O–C(O)–, indanyl$(CH_2)_{0-3}$–O–C(O)–, aryl$(CH_2)_{1-3}$–C(O)–, aryl–cyclopropyl–C(O)–, heteroaryl–cyclopropyl–C(O)–, heteroaryl$(CH_2)_{1-3}$–C(O)–, aryl$(CH_2)_{1-3}$–, heteroaryl$(CH_2)_{1-3}$–, aryl$(CH_2)_{1-3}$–NH–C(O)–, aryl$(CH_2)_{1-3}$–NH–C(NCN)–, aryl$(CH_2)_{1-3}$–SO_2–, heteroaryl$(CH_2)_{1-3}$–SO_2–, wherein any of the aryl or heteroaryl is optionally substituted by 1-5 substitutents, each substituent independently is C_{1-4}alkyl, C_{3-6}cycloalkyl, C_{1-4}alkoxy, trifluoromethyl, bromo, fluoro, or chloro; and

X is H, OH, F, C_{1-4}alkyl, C_{1-4}alkoxy, NH_2, or X taken with an adjacent bond is =O.

2. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

B is aryl$(CH_2)_{0-3}$–O–C(O)–, wherein the aryl is optionally substituted by 1-5 substitutents, each substituent independently is C_{1-4}alkyl, C_{3-6}cycloalkyl, C_{1-4}alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

3. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 1 nitrogen ring atom;

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

4. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is an isoxazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

5. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is a thiadiazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-}

4alkyl)(C_0-4alkyl), nitro, (C_1-2alkyl)(C_1-2alkyl)NCH_2–, (C_1-2alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

6. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 5 membered heteroaromatic ring containing 2 nitrogen ring atoms;

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C_1-4alkyl, C_1-4alkoxy, C_2-4alkynyl, trifluoromethyl, hydroxy, hydroxyC_1-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_0-4alkyl)(C_0-4alkyl), nitro, (C_1-2alkyl)(C_1-2alkyl)NCH_2–, (C_1-2alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

7. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is quinolinyl optionally substituted with 1 or 2 substituents, each substituent independently is C_1-4alkyl, C_1-4alkoxy, C_2-4alkynyl, trifluoromethyl, hydroxy, hydroxyC_1-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_0-4alkyl)(C_0-4alkyl), nitro, (C_1-2alkyl)(C_1-2alkyl)NCH_2–, (C_1-2alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

8. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is purinyl optionally substituted with 1 or 2 substituents, each substituent independently is C_1-4alkyl, C_1-4alkoxy, C_2-4alkynyl, trifluoromethyl, hydroxy, hydroxyC_1-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_0-4alkyl)(C_0-4alkyl), nitro, (C_1-2alkyl)(C_1-2alkyl)NCH_2–, (C_1-2alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

9. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 2 nitrogen ring atoms;

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

10. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is thiazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

11. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is pteridinyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

12. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is pyrrolopyrimidinyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

13. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is a imidazopyridinyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

14. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is benzimidazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

15. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

B is aryl$(CH_2)_{1-3}$–SO_2–, wherein the aryl is optionally substituted by 1-5 substitutents, each substituent independently is C_{1-4}alkyl, C_{3-6}cycloalkyl, C_{1-4}alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

16. The compound according to Claim 15, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 2 nitrogen ring atoms;

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl),

nitro, $(C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–$, $(C_{1-2}alkyl)HNCH_2–$, $Si(CH_3)_3–C–$, or $NH_2C(O)–$.

17. The compound according to Claim 15, or pharmaceutically acceptable salts thereof, wherein

HetAr is quinazolinyl optionally substituted with 1 or 2 substituents, each substituent independently is $C_{1-4}alkyl$, $C_{1-4}alkoxy$, $C_{2-4}alkynyl$, trifluoromethyl, hydroxy, hydroxy$C_{1-4}alkyl$, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–$N(C_{0-4}alkyl)(C_{0-4}alkyl)$, nitro, $(C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–$, $(C_{1-2}alkyl)HNCH_2–$, $Si(CH_3)_3–C–$, or $NH_2C(O)–$.

18. The compound according to Claim 15, or pharmaceutically acceptable salts thereof, wherein

HetAr is purinyl optionally substituted with 1 or 2 substituents, each substituent independently is $C_{1-4}alkyl$, $C_{1-4}alkoxy$, $C_{2-4}alkynyl$, trifluoromethyl, hydroxy, hydroxy$C_{1-4}alkyl$, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–$N(C_{0-4}alkyl)(C_{0-4}alkyl)$, nitro, $(C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–$, $(C_{1-2}alkyl)HNCH_2–$, $Si(CH_3)_3–C–$, or $NH_2C(O)–$.

19. The compound according to Claim 15, or pharmaceutically acceptable salts thereof, wherein

HetAr is imidazopyridinyl optionally substituted with 1 or 2 substituents, each substituent independently is $C_{1-4}alkyl$, $C_{1-4}alkoxy$, $C_{2-4}alkynyl$, trifluoromethyl, hydroxy, hydroxy$C_{1-4}alkyl$, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–$N(C_{0-4}alkyl)(C_{0-4}alkyl)$, nitro, $(C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–$, $(C_{1-2}alkyl)HNCH_2–$, $Si(CH_3)_3–C–$, or $NH_2C(O)–$.

20. The compound according to Claim 15, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 1 nitrogen ring atom; and

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)HNCH_2–, Si$(CH_3)_3$–C–, or $NH_2C(O)$–.

21. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 5 membered ring containing 1 nitrogen ring atom; and

B is aryl$(CH_2)_{0-3}$–O–C(O)–, wherein the aryl is optionally substituted by 1-5 substitutents, each substituent independently is C_{1-4}alkyl, C_{3-6}cycloalkyl, C_{1-4}alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

22. The compound according to Claim 21, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 2 nitrogen ring atoms;

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)HNCH_2–, Si$(CH_3)_3$–C–, or $NH_2C(O)$–.

23. The compound according to Claim 21, or pharmaceutically acceptable salts thereof, wherein

HetAr is pteridinyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)HNCH_2–, Si$(CH_3)_3$–C–, or $NH_2C(O)$–.

24. The compound according to Claim 21, or pharmaceutically acceptable salts thereof, wherein

HetAr is purinyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

25. The compound according to Claim 21, or pharmaceutically acceptable salts thereof, wherein

HetAr is benzimidazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

26. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is an aza bicyclo octane ring; and

B is aryl$(CH_2)_{0-3}$–O–C(O)–, wherein the aryl is optionally substituted by 1-5 substitutents, each substituent independently is C_{1-4}alkyl, C_{3-6}cycloalkyl, C_{1-4}alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

27. The compound according to Claim 26, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 1 nitrogen ring atom; and

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

28. The compound according to Claim 26, or pharmaceutically acceptable salts thereof, wherein

HetAr is purinyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)HNCH_2–, Si$(CH_3)_3$–C–, or $NH_2C(O)$–.

29. The compound according to Claim 26, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 2 nitrogen ring atom; and

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)HNCH_2–, Si$(CH_3)_3$–C–, or $NH_2C(O)$–.

30. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is an aza bicyclo octane ring; and

B is aryl$(CH_2)_{1-3}$–SO_2–, wherein the aryl is optionally substituted by 1-5 substitutents, each substituent independently is C_{1-4}alkyl, C_{3-6}cycloalkyl, C_{1-4}alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

31. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

B is heteroaryl$(CH_2)_{1-3}$–C(O)–, wherein the heteroaryl is optionally substituted by 1-5 substitutents, each substituent independently is C_{1-4}alkyl, C_{3-6}cycloalkyl, C_{1-4}alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

32. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

B is aryl$(CH_2)_{1-3}$–C(O)–, wherein the aryl is optionally substituted by 1-5 substitutents, each substituent independently is C_{1-4}alkyl, C_{3-6}cycloalkyl, C_{1-4}alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

33. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

B is aryl–cyclopropyl–C(O)–, wherein the aryl is optionally substituted by 1-5 substitutents, each substituent independently is C_{1-4}alkyl, C_{3-6}cycloalkyl, C_{1-4}alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

34. The compound according to Claim 33, or pharmaceutically acceptable salts thereof, wherein

HetAr is pyridyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

35. The compound according to Claim 33, or pharmaceutically acceptable salts thereof, wherein

HetAr is pyrazinyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

36. The compound according to Claim 33, or pharmaceutically acceptable salts thereof, wherein

HetAr is pyridazinyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

37. The compound according to Claim 33, or pharmaceutically acceptable salts thereof, wherein

HetAr is pyrimidinyl optionally substituted with 1 or 2 substituents, each substituent independently is C_{1-4}alkyl, C_{1-4}alkoxy, C_{2-4}alkynyl, trifluoromethyl, hydroxy, hydroxyC_{1-4}alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl–, phenylethynyl–, heteroarylethynyl–,–N(C_{0-4}alkyl)(C_{0-4}alkyl), nitro, (C_{1-2}alkyl)(C_{1-2}alkyl)NCH_2–, (C_{1-2}alkyl)$HNCH_2$–, $Si(CH_3)_3$–C–, or $NH_2C(O)$–.

38. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

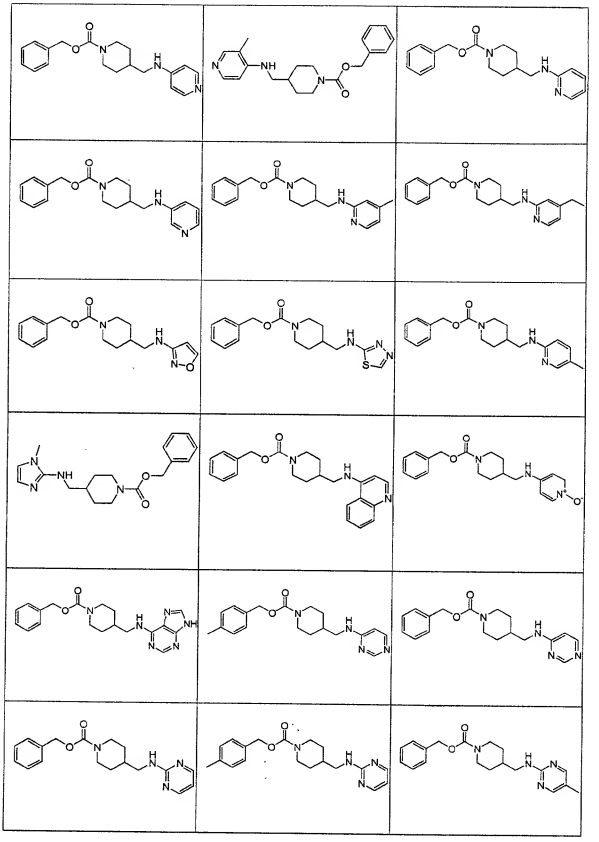
B is heteroaryl$(CH_2)_{1-3}$–O–C(O)–, wherein the heteroaryl is optionally substituted by 1-5 substitutents, each substituent independently is C_{1-4}alkyl, C_{3-6}cycloalkyl, C_{1-4}alkoxy, trifluoromethyl, bromo, fluoro, or chloro;.

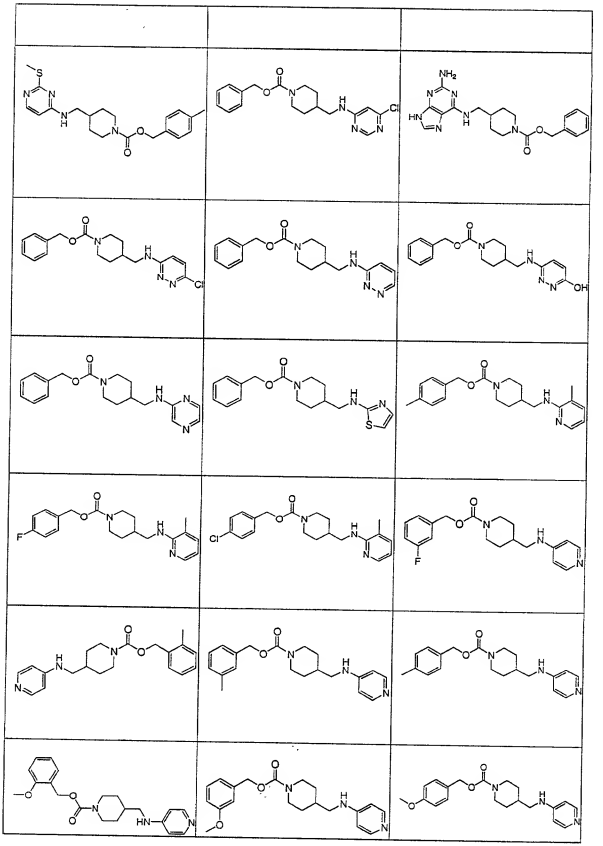
39. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

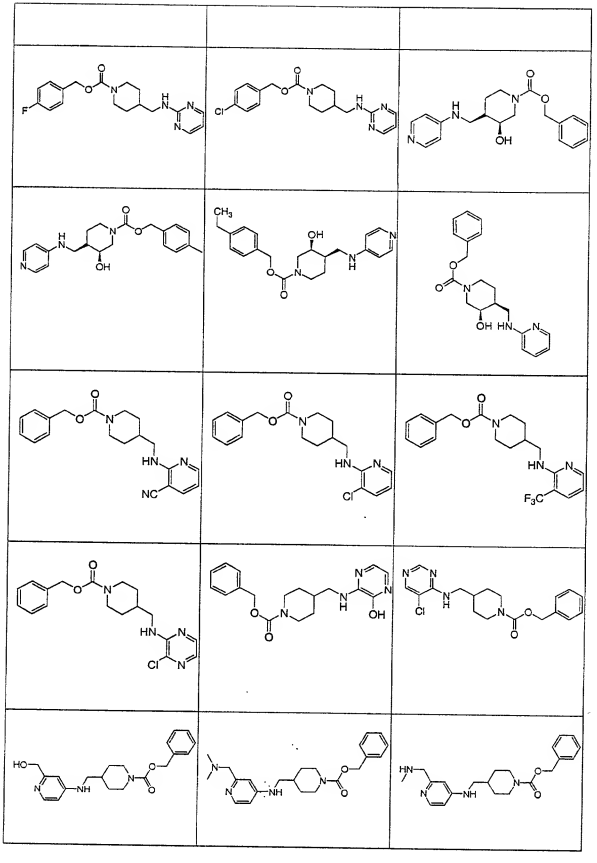
NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

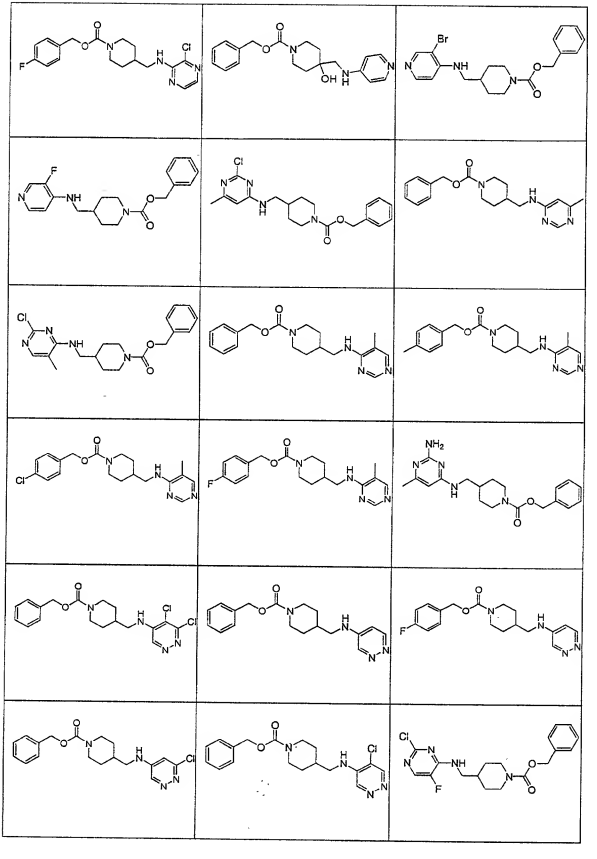
B is aryl$(CH_2)_{1-3}$–NH–C(NCN)–, wherein the aryl is optionally substituted by 1-5 substitutents, each substituent independently is C_{1-4}alkyl, C_{3-6}cycloalkyl, C_{1-4}alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

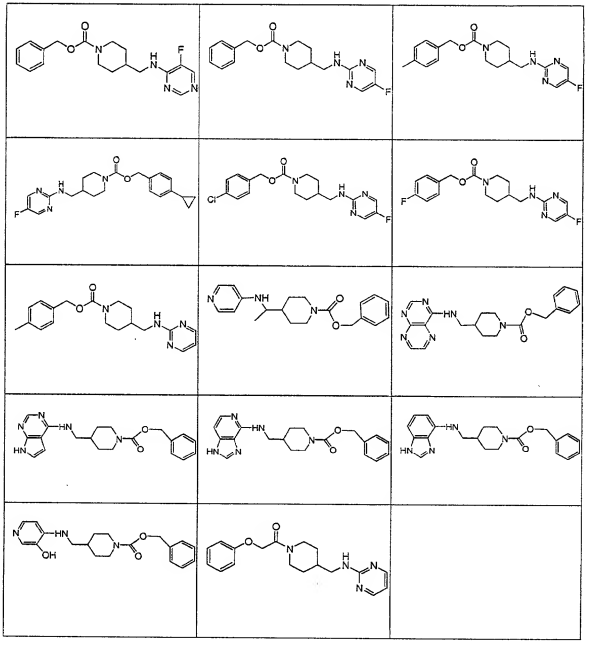
40. The compound according to Claim 1, wherein said compound is





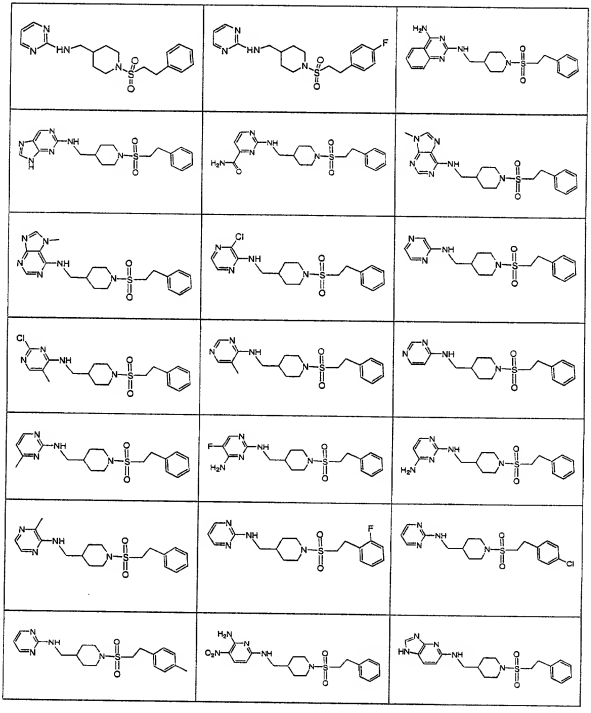






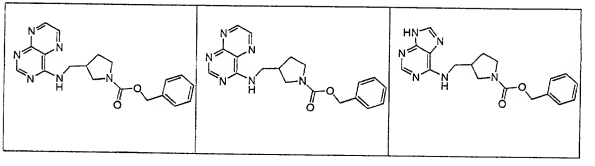
or a pharmaceutically acceptable salt thereof.

41. The compound according to Claim 1, wherein said compound is



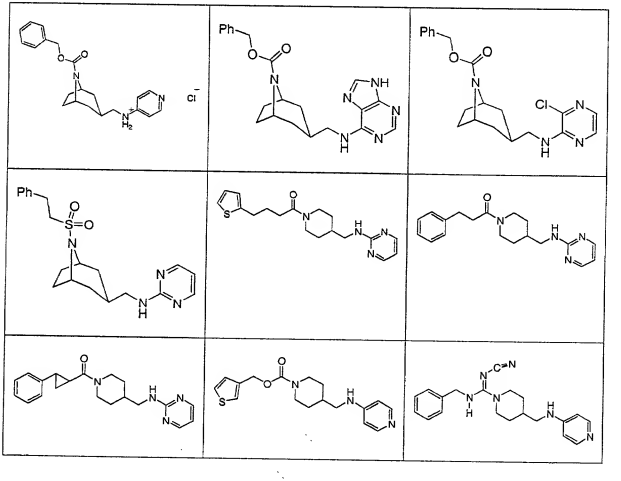
or a pharmaceutically acceptable salt thereof.

42. The compound according to Claim 1, wherein said compound is



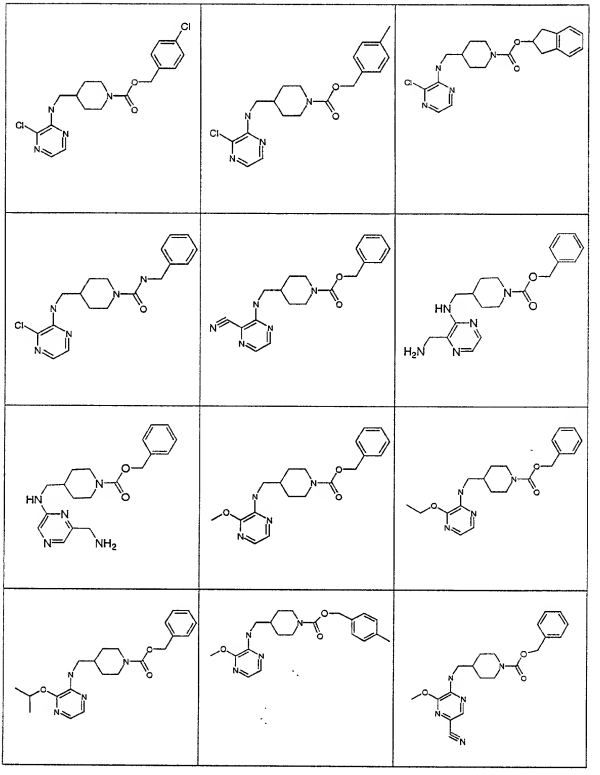
or a pharmaceutically acceptable salt thereof.

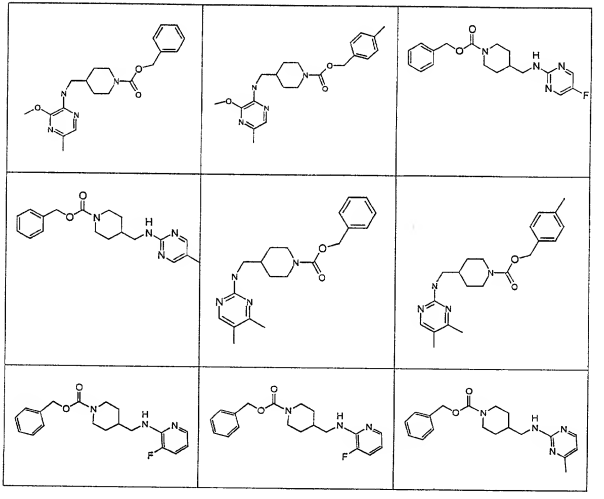
43. The compound according to Claim 1, wherein said compound is



or a pharmaceutically acceptable salt thereof.

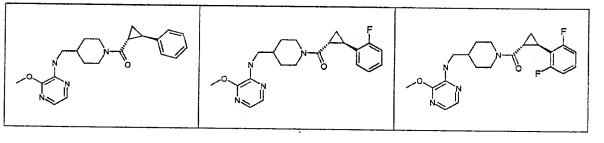
44. The compound according to Claim 1, wherein said compound is

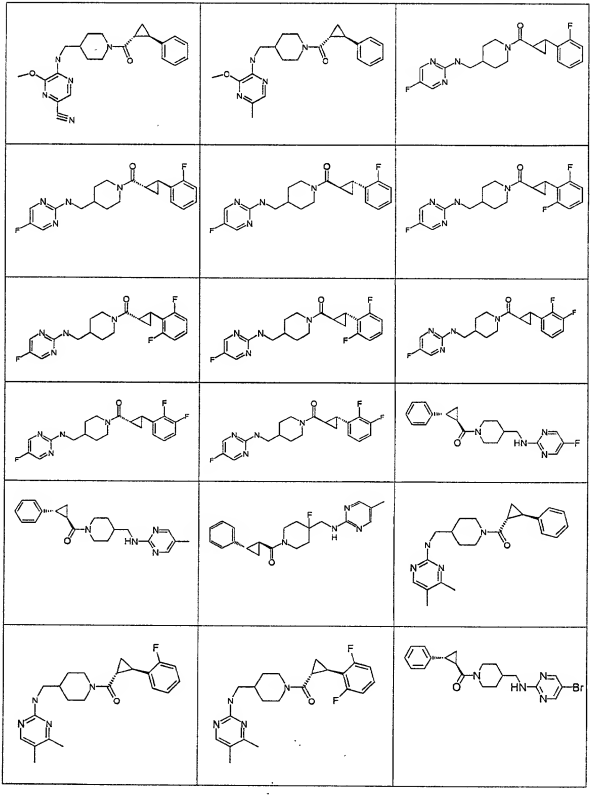


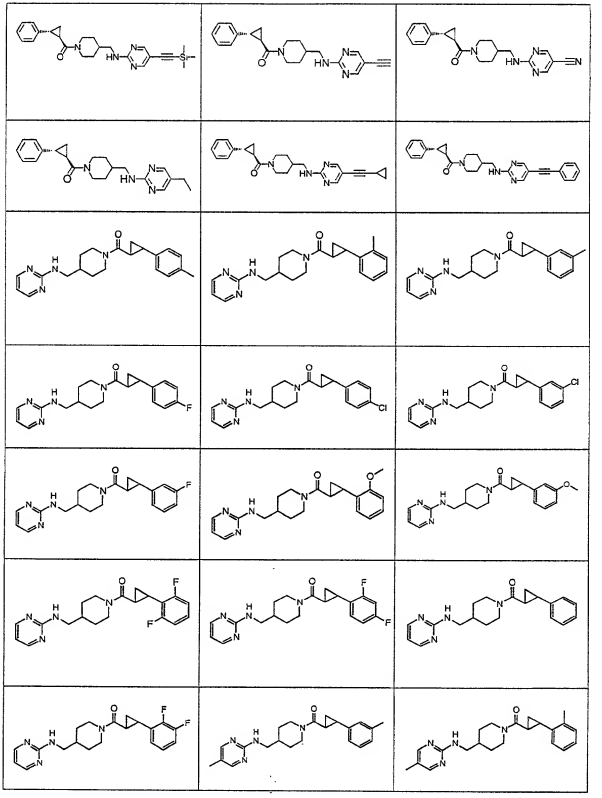


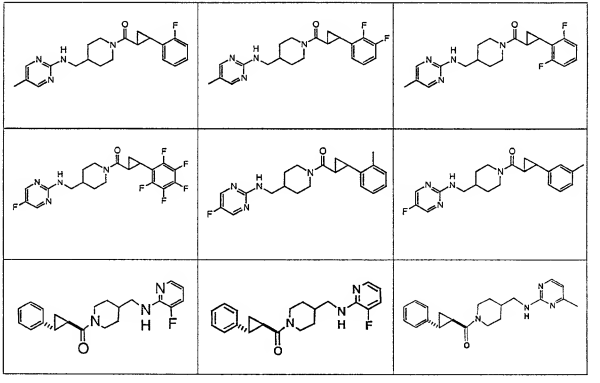
or a pharmaceutically acceptable salt thereof.

45. The compound according to Claim 1, wherein said compound is



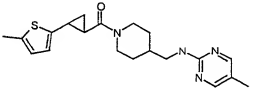






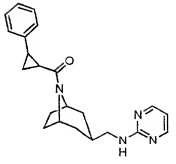
or a pharmaceutically acceptable salt thereof.

46. The compound according to Claim 1, wherein said compound is



or a pharmaceutically acceptable salt thereof.

47. The compound according to Claim 1, wherein said compound is



or a pharmaceutically acceptable salt thereof.

48. A pharmaceutical composition comprising an inert carrier and an effective amount of a compound according to claim 1.

49. The pharmaceutical composition according to claim 48 useful for the treatment of pain.

50. The pharmaceutical composition according to claim 48 useful for the treatment of migraine, depression, anxiety, schizophrenia, Parkinson's disease, or stroke.

51. A method of treating pain comprising a step of administering to one in need of such treatment an effective amount of a compound according to claim 1.

52. A method of treating migraine, depression, anxiety, schizophrenia, Parkinson's disease, or stroke comprising a step of administering to one in need of such treatment an effective amount of a compound according to claim 1.